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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO
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7590 01/26/2004			EXAMINER	
CARELLA, BYRNE, BAIN, GILFILLAN, CECCHI, STEWART & OLSTEIN			SMITH, CAROLYN L	
6 Becker Farm Road			ART UNIT	PAPER NUMBER
Roseland, NJ	07068		1631	
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Please find below and/or attached an Office communication concerning this application or proceeding.

-	Application No.	Applicant(s) WEAVER, ZOE	
	09/954,531		
Office Action Summary	Examiner	Art Unit	
	Carolyn L Smith	1631	
The MAILING DATE of this communication Period for Reply	appears on the cover sheet w	ith the correspondence address	
A SHORTENED STATUTORY PERIOD FOR RETHE MAILING DATE OF THIS COMMUNICATION - Extensions of time may be available under the provisions of 37 CF after SIX (6) MONTHS from the mailing date of this communication - If the period for reply specified above is less than thirty (30) days, and the period for reply is specified above, the maximum statutory period for reply within the set or extended period for reply will, by second and the period for reply will, by second period for reply will, by second period for reply will, by second patent term adjustment. See 37 CFR 1.704(b).	DN. R 1.136(a). In no event, however, may a in the statutory minimum of this a reply within the statutory minimum of this eriod will apply and will expire SIX (6) MON tatute, cause the application to become Af	reply be timely filed ty (30) days will be considered timely. ITHS from the mailing date of this communication. BANDONED (35 U.S.C. § 133).	
1) Responsive to communication(s) filed on 1	0 November 2003.		
, ·	his action is non-final.		
Since this application is in condition for all closed in accordance with the practice und			
Disposition of Claims			
4) ☐ Claim(s) <u>1-35 and 37-48</u> is/are pending in 4a) Of the above claim(s) <u>18-35 and 37-46</u> 5) ☐ Claim(s) is/are allowed. 6) ☐ Claim(s) <u>1-17,47 and 48</u> is/are rejected. 7) ☐ Claim(s) <u>1, 3, 6-17, 47, and 48</u> is/are objection is/are objection is/are subject to res	is/are withdrawn from conside		
Application Papers			
9) The specification is objected to by the Exam 10) The drawing(s) filed on is/are: a) Applicant may not request that any objection to Replacement drawing sheet(s) including the co 11) The oath or declaration is objected to by the	accepted or b) objected to the drawing(s) be held in abeyar rrection is required if the drawing	nce. See 37 CFR 1.85(a). (s) is objected to. See 37 CFR 1.121(d).	
Priority under 35 U.S.C. §§ 119 and 120			
12) Acknowledgment is made of a claim for for a) All b) Some * c) None of: 1. Certified copies of the priority docum 2. Certified copies of the priority docum 3. Copies of the certified copies of the papplication from the International Bu * See the attached detailed Office action for a 13) Acknowledgment is made of a claim for dom since a specific reference was included in the 37 CFR 1.78. a) The translation of the foreign language 14) Acknowledgment is made of a claim for dom reference was included in the first sentence of	nents have been received. The nents have been received in A priority documents have been reau (PCT Rule 17.2(a)). The list of the certified copies not restic priority under 35 U.S.C. of first sentence of the specifical provisional application has beestic priority under 35 U.S.C.	pplication No received in this National Stage received. § 119(e) (to a provisional application ation or in an Application Data Sheet een received. §§ 120 and/or 121 since a specific	
Attachment(s)			
Notice of References Cited (PTO-892) Notice of Draftsperson's Patent Drawing Review (PTO-948) Information Disclosure Statement(s) (PTO-1449) Paper No.) 5) Notice of h	Summary (PTO-413) Paper No(s) nformal Patent Application (PTO-152)	

DETAILED ACTION

Applicant's amendments and remarks, filed 11/10/03, are acknowledged. Amended claims 1, 2, 11, 14, 15, 47, and 48 are acknowledged. On page 9 (first sentence) of Applicant's response, it is incorrectly noted that claims 1-17, 47, and 48 are pending. Actually, claims 1-35 and 37-48 are pending while claims 18-35 and 37-46 are withdrawn from consideration as being drawn to non-elected Groups.

Applicant's arguments, filed 11/10/03, have been fully considered but they are not deemed to be persuasive. Rejections and/or objections not reiterated from the previous office actions are hereby withdrawn. The following rejections and/or objections are either reiterated or newly applied. They constitute the complete set presently being applied to the instant application.

Claims 1-17, 47, and 48 are herein under examination.

Claim Objections

Claims 1, 3, 6-17, 47, and 48 are objected to due to their inclusion of subject matter that has been non-elected due to a restriction requirement and therefore withdrawn from consideration. The non-elected subject matter of claims 1, 3, 6-17, 47, and 48 is as follows: Claims 1, 3, 6-17, 47, and 48 contain sequences, such as sequences other than SEQ ID NO: 110,

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653, 683, 767, 804, 820, 910, 1019, 1040, and 1247, which are non-elected subject matter. Removal of non-elected subject matter is requested.

Claim Rejections - 35 U.S.C. 112, first paragraph

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

LACK OF WRITTEN DESCRIPTION

Claims 1-17, 47, and 48 are rejected under 35 U.S.C. 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor, at the time of the invention was filed, had possession of the claimed invention.

The rejection of claims 1-17, 47, and 48 is maintained under 35 U.S.C. 112 as containing subject matter which was not described in the specification in such a way as to reasonably convey to one skilled in the relevant art that the inventor, at the time of the invention was filed, had possession of the claimed invention.

The specification discloses SEQ ID NO: 110, 653, 683, 767, 804, 820, 910, 1019, 1040, and 1247 which correspond to nucleic acid sequences. SEQ ID NO: 110, 653, 683, 767, 804, 820, 910, 1019, 1040, and 1247 and their full complements meet the written description provisions of 35 U.S.C. 112, first paragraph. However, due to the open claim language of "containing a gene that corresponds to a polynucleotide" (claim 1) and "comprising a nucleotide sequence corresponding to a gene" (claim 48), these claims encompass sequences which do not

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meet the written description provision of 35 U.S.C. 112, first paragraph. The specification provides insufficient written description to support the genus encompassed by these claims.

<u>Vas-Cath Inc. v. Mahurkar</u>, 19 USPQ2d 1111, makes clear that "applicant must convey with reasonable clarity to those skilled in the art that, as of the filing date sought, he or she was in possession *of the invention*. The invention is, for purposes of the 'written description' inquiry, whatever is now claimed." (See page 1117.) The specification does not "clearly allow persons of ordinary skill in the art to recognize that [he or she] invented what is claimed." (See <u>Vas-Cath</u> at page 1116.)

With the exception of SEQ ID NO: 110, 653, 683, 767, 804, 820, 910, 1019, 1040, and 1247, the skilled artisan cannot envision the detailed chemical structure of the encompassed polynucleotides and/or proteins, regardless of the complexity or simplicity of the method of isolation. Adequate written description requires more than a mere statement that it is part of the invention and reference to a potential method for isolating it. The nucleic acid itself is required. See Fiers v. Revel, 25 USPQ2d 1601, 1606 (CAFC 1993) and Amgen Inc. V. Chugai Pharmacentical Co. Ltd., 18 USPQ2d 1016. In Fiddes v. Baird, 30 USPQ2d 1481, 1483, claims directed to mammalian FGF's were found unpatentable due to lack of written description for the broad class. The specification provided only the bovine sequence.

Finally, <u>University of California v. Eli Lilly and Co.</u>, 43 USPQ2d 1398, 1404, 1405 held that:

...To fulfill the written description requirement, a patent specification must describe an invention and do so in sufficient detail that one skilled in the art can clearly conclude that "the inventor invented the claimed invention." *Lockwood v. American Airlines, Inc.*, 107 F.3d 1565, 1572, 41 USPQ2d 1961, 1966 (1997); *In re Gosteli*, 872 F.2d 1008, 1012, 10 USPQ2d 1614, 1618 (Fed. Cir. 1989) (" [T]he description must clearly allow persons of ordinary skill in the art to recognize that [the inventor] invented what is claimed."). Thus, an applicant complies with the written description requirement "by describing the invention, with all its claimed limitations, not that which makes it obvious," and by using "such descriptive means as words, structures, figures, diagrams, formulas, etc., that set forth the claimed invention." *Lockwood*, 107 F.3d at 1572, 41 USPQ2d at 1966.

Therefore, only SEQ ID NO: 110, 653, 683, 767, 804, 820, 910, 1019, 1040, and 1247 and their full-length complements, but not the full breadth of the claims meet the written description provision of 35 USC 112, first paragraph. The species specifically disclosed are not representative of the genus because the genus is highly variant. Applicant is reminded that <u>Vas-</u>

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Cath makes clear that the written description provision of 35 USC 112 is severable from its

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enablement provision. (See page 1115.)

Applicant correctly notes that claim 48, not claim 54, was rejected for failing to meet the written description requirements for the phrase "comprising a nucleotide sequence corresponding to a gene". Applicant states this rejection (as well as the phrase "containing a gene that corresponds to a polynucleotide" in claim 1), cannot be solely based on the use of open ended language because "comprising" is standard claim language and that this rejection must be predicated on the use of the term "corresponding" in conjunction with "comprising". It is noted that this rejection is made because these phrases, in their broadest and reasonable interpretation. encompass sequences that do not have written support in the specification, claims, and/or drawings as originally filed. Applicant states the term "correspond" is defined as meaning a gene that encodes an RNA at least 90% identical to the claimed polynucleotide. This is found unpersuasive as this sequence could encompass a claimed sequence, plus up to 10% of additional sequence on either end that does not meet the written description provision of 35 U.S.C. 112, first paragraph. Again, it is acknowledged that the Applicant has written support for SEO ID NO: 110, 653, 683, 767, 804, 820, 910, 1019, 1040, and 1247 and their full length complements. but not for the full breath of the claims. The arguments presented by the Applicant do not specifically address the written description rejection, as reiterated, and is therefore found unpersuasive.

Claim Rejections - 35 U.S.C. 112, first paragraph

The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or

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with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Factors to be considered in determining whether a disclosure would require undue experimentation have been summarized in *Ex parte Forman*, 230 USPQ 546 (BPAI 1986) and reiterated by the Court of Appeals in *In re Wands*, 8 USPQ2d 1400 at 1404 (CAFC 1988). The factors to be considered in determining whether undue experimentation is required include: (1) the quantity of experimentation necessary, (2) the amount or direction presented, (3) the presence or absence of working examples, (4) the nature of the invention, (5) the state of the prior art, (6) the relative skill of those in the art, (7) the predictability or unpredictability of the art, and (8) the breadth of the claims. The Board also stated that although the level of the skill in molecular biology is high, the results of experiments in genetic engineering are unpredictable. While all of these factors are considered, a sufficient amount for a *prima facie* case are discussed below.

LACK OF ENABLEMENT

Claims 1-17, 47, and 48 are rejected under 35 U.S.C. § 112, first paragraph, as containing subject matter which was not described in the specification in such a way as to enable one skilled in the art to which it pertains, or with which it is most nearly connected, to make and/or use the claimed invention.

The sequences selected in these methods include sequences from GenBank with accession numbers AP001082, AA478962, N64489, AA227221, D80055, N45300, D60118, H02533, AA010665, and N69022. AP001082 is from genomic human DNA from chromosome 11; AA478962 is from cDNA from pooled human melanocyte, fetal heart, and pregnant uterus; N64489 is from cDNA from adult male multiple sclerosis lesions tissue; AA227221 is from cDNA from human brain neuroepithelial cells; D80055 is from cDNA from human fetal brain;

N45300 is from cDNA from adult male multiple sclerosis lesions tissue; D60118 is from cDNA from human fetal brain; H02533 is from cDNA from human female placenta; AA010665 is from cDNA from human fetal heart; and N69022 is from cDNA from human fetal lung. There are millions of sequences in the world with a small portion actually available in public databases, such as GenBank. A microarray type of invention that involves a multitude of sequences that appear to be randomly selected with no previously known function or association with cancer that merely come from chromosome 11, melanocytes, fetal organs, female organs, multiple sclerosis lesions, brain neuroepithelial cells (as is the case with the elected sequences) do not appear to be enabling for screening chemical compounds for anti-neoplastic activity. The quantity of experimentation required to verify that these sequences represent valid predictors of screening chemical compounds for anti-neoplastic activity appears to be undue. Due to undue experimentation required, the lack of guidance directed to verifying such sequences functioning as valid predictors, the lack of working examples addressing the same, the unpredictability of knowing if these sequences are potentially valid predictors for screening anti-neoplastic activity. and the breath of the claims; this invention is rejected due to the lack of enablement for one skilled in the art to be able to make and use the invention.

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Conclusion

No claim is allowed.

Papers related to this application may be submitted to Technical Center 1600 by facsimile transmission. Papers should be faxed to Technical Center 1600 via the PTO Fax Center located

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in Crystal Mall 1. The faxing of such papers must conform with the notices published in the Official Gazette, 1096 OG 30 (November 15, 1988), 1156 OG 61 (November 16, 1993), and 1157 OG 94 (December 28, 1993) (See 37 CFR §1.6(d)). The CM1 Fax Center number is (703) 872-9306.

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Carolyn Smith, whose telephone number is (571) 272-0721. The examiner can normally be reached Monday through Thursday from 8 A.M. to 6:30 P.M.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Michael Woodward, can be reached on (571) 272-0722.

Any inquiry of a general nature or relating to the status of this application should be directed to Legal Instruments Examiner Tina Plunkett whose telephone number is (571) 272-0549 or to the Technical Center receptionist whose telephone number is (703) 308-0196.

January 13, 2004

ARDIN H. MARSCHEL PRIMARY EXAMELER